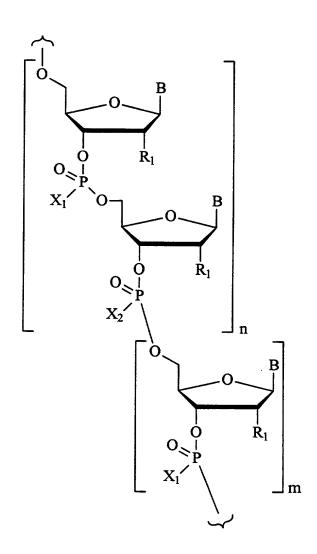
WHAT IS CLAIMED IS:

1. A compound comprising a plurality of covalently-bound 2'-modified nucleosides having the formula:



5 wherein:

each B is a nucleobase;

one of X₁ or X₂ is O, and the other of X₁ or X₂ is S; each R₁, is, independently, H, hydroxyl, C₁-C₂₀ alkyl, C₃-C₂₀ alkenyl, C₂-C₂₀ alkynyl, halogen, thiol, keto, carboxyl, nitro, nitroso, nitrile, trifluoromethyl, trifluoromethoxy, O-alkyl, S-alkyl, NH-alkyl, N-dialkyl, O-aryl, S-aryl, NH-aryl, O-aralkyl, S-aralkyl, NH-aralkyl, amino, N-

phthalimido, imidazole, azido, hydrazino, hydroxylamino,
isocyanato, sulfoxide, sulfone, sulfide, disulfide, silyl,
aryl, heterocycle, carbocycle, intercalator, reporter
molecule, conjugate, polyamine, polyamide, polyalkylene
5 glycol, or polyether;

or R_1 is a group of formula $Z-R_{22}-(R_{23})_v$; $Z \text{ is C, S, NH, or N-}R_{22}-(R_{23})_v;$ $R_{22} \text{ is } C_1-C_{20} \text{ alkyl, } C_2-C_{20} \text{ alkenyl, or } C_2-C_{20}$ alkynyl;

10 R₂₃ is hydrogen, amino, halogen, hydroxyl, thiol, keto, carboxyl, nitro, nitroso, nitrile, trifluoromethyl, trifluoromethoxy, O-alkyl, S-alkyl, NH-alkyl, N-dialkyl, O-aryl, S-aryl, NH-aryl, O-aralkyl, S-aralkyl, NH-aralkyl, amino, N-phthalimido, imidazole, azido, hydrazino, hydroxylamino, isocyanato, sulfoxide, sulfone, sulfide, disulfide, silyl, aryl, heterocycle, carbocycle, intercalator, reporter molecule, conjugate, polyamine, polyamide, polyalkylene glycol, polyether, a group that enhances the pharmacodynamic properties of oligonucleotides, or a group

20 that enhances the pharmacokinetic properties of oligonucleotides;

v is from 0 to about 10; or R_1 has the formula:

$$-(O)_{y_1} - (CH_2)_{y_2} - O - N - (CH_2)_{y_2} - O - E$$

y1 is 0 or 1; y2 is independently 0 to 10; y3 is 1 to 10; E is C_1-C_{10} alkyl, $N(Q_1)(Q_2)$ or $N=C(Q_1)(Q_2)$; each Q_1 and Q_2 is, independently, H, C_1 - C_{10} alkyl, substituted alkyl, dialkylaminoalkyl, a nitrogen protecting group, a tethered or untethered conjugate group, a linker to a solid support; or Q_1 and Q_2 , together, are joined in a nitrogen protecting group or a ring structure that can include at least one additional heteroatom selected from N and O;

or R_1 has one of formula I or II:

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wherein

 Z_0 is O, S, or NH;

 q^1 is from 0 to 10;

 q^2 is from 1 to 10;

 q^3 is 0 or 1;

q4 is, 0, 1 or 2;

 Z_4 is OM_1 , SM_1 , or $N(M_1)_2$;

each M_1 is, independently, H, C_1-C_8 alkyl, C_1-C_8

20 haloalkyl, $C(=NH)N(H)M_2$, $C(=O)N(H)M_2$ or $OC(=O)N(H)M_2$;

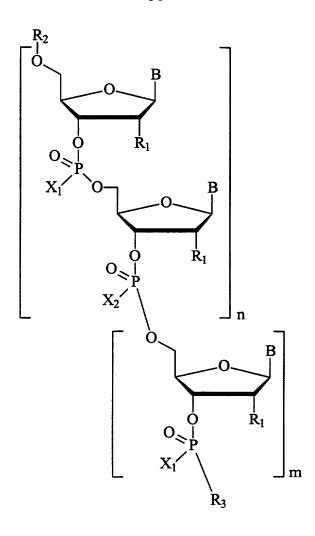
 M_2 is H or C_1-C_8 alkyl;

 Z_1 , Z_2 and Z_3 comprise a ring system having from about 4 to about 7 carbon atoms, or having from about 3 to about 6 carbon atoms and 1 or 2 hetero atoms wherein said 25 hetero atoms are selected from oxygen, nitrogen and sulfur, and wherein said ring system is aliphatic, unsaturated aliphatic, aromatic, or saturated or unsaturated heterocyclic; and

 Z_5 is alkyl or haloalkyl having 1 to about 10 carbon atoms, alkenyl having 2 to about 10 carbon atoms, alkynyl having 2 to about 10 carbon atoms, aryl having 6 to about 14 carbon atoms, $N\left(Q_1\right)\left(Q_2\right)$, OQ_1 , halo, SQ_1 or CN;

5 n is from 2 to 50; and m is 0 or 1.

- 2. The compound of claim 1 wherein R_1 is $-\text{O-CH}_2\text{-CH}_2\text{-O-CH}_3$.
- 3. The compound of claim 1 wherein n is about 5 to 10 about 50.
 - 4. The compound of claim 1 wherein n is about 8 to about 30.
 - 5. The compound of claim 1 wherein n is about 4 to about 15.
- 15 6. The compound of claim 1 wherein n is 2 to about 10.
 - 7. An oligonucleotide having the Formula:



wherein:

each B is a nucleobase;

 X_1 is S;

5 X_2 is 0;

each R₁, is, independently, H, hydroxyl, C₁-C₂₀ alkyl, C₃-C₂₀ alkenyl, C₂-C₂₀ alkynyl, halogen, thiol, keto, carboxyl, nitro, nitroso, nitrile, trifluoromethyl, trifluoromethoxy, O-alkyl, S-alkyl, NH-alkyl, N-dialkyl, O-aryl, S-aryl, NH-aryl, O-aralkyl, S-aralkyl, NH-aralkyl, amino, N-phthalimido, imidazole, azido, hydrazino, hydroxylamino, isocyanato, sulfoxide, sulfone, sulfide, disulfide, silyl, aryl, heterocycle, carbocycle, intercalator, reporter

molecule, conjugate, polyamine, polyamide, polyalkylene glycol, or polyether;

or R_1 is a group of formula $Z-R_{22}-(R_{23})_v$;

Z is O, S, NH, or $N-R_{22}-(R_{23})_{y}$;

 R_{22} is C_1-C_{20} alkyl, C_2-C_{20} alkenyl, or C_2-C_{20}

alkynyl;

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R₂₃ is hydrogen, amino, halogen, hydroxyl, thiol, keto, carboxyl, nitro, nitroso, nitrile, trifluoromethyl, trifluoromethoxy, O-alkyl, S-alkyl, NH-alkyl, N-dialkyl, O-aryl, S-aryl, NH-aryl, O-aralkyl, S-aralkyl, NH-aralkyl, amino, N-phthalimido, imidazole, azido, hydrazino, hydroxylamino, isocyanato, sulfoxide, sulfone, sulfide, disulfide, silyl, aryl, heterocycle, carbocycle, intercalator, reporter molecule, conjugate, polyamine, polyamide, polyalkylene glycol, polyether, a group that enhances the pharmacodynamic properties of oligonucleotides, or a group that enhances the pharmacokinetic properties of oligonucleotides;

v is from 0 to about 10; or R_1 has the formula:

$$-(O)_{y_1} - (CH_2)_{y_2} - O - N - (CH_2)_{y_2} - O - E$$

y1 is 0 or 1;

y2 is independently 0 to 10;

y3 is 1 to 10;

E is C_1-C_{10} alkyl, $N\left(Q_1\right)\left(Q_2\right)$ or $N=C\left(Q_1\right)\left(Q_2\right)$;

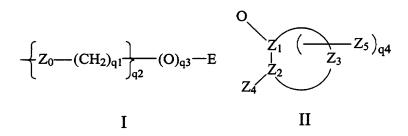
each Q_1 and Q_2 is, independently, H, C_1 - C_{10} alkyl, substituted alkyl, dialkylaminoalkyl, a nitrogen protecting group, a tethered or untethered conjugate group,

a linker to a solid support; or Q_1 and Q_2 , together, are joined in a nitrogen protecting group or a ring structure that can include at least one additional heteroatom selected from N and O;

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or R_1 has one of formula I or II:



wherein

 Z_0 is O, S, or NH;

 q^1 is from 0 to 10;

 q^2 is from 1 to 10;

 q^3 is 0 or 1;

q4 is, 0, 1 or 2;

15 Z_4 is OM_1 , SM_1 , or $N(M_1)_2$;

each M_1 is, independently, H, C_1-C_8 alkyl, C_1-C_8 haloalkyl, $C(=NH)\,N\,(H)\,M_2$, $C\,(=O)\,N\,(H)\,M_2$ or $OC\,(=O)\,N\,(H)\,M_2$;

 M_2 is H or C_1-C_8 alkyl;

 Z_1 , Z_2 and Z_3 comprise a ring system having from 20 about 4 to about 7 carbon atoms, or having from about 3 to about 6 carbon atoms and 1 or 2 hetero atoms wherein said hetero atoms are selected from oxygen, nitrogen and sulfur, and wherein said ring system is aliphatic, unsaturated aliphatic, aromatic, or saturated or unsaturated

25 heterocyclic; and

 Z_{5} is alkyl or haloalkyl having 1 to about 10 carbon atoms, alkenyl having 2 to about 10 carbon atoms,

alkynyl having 2 to about 10 carbon atoms, aryl having 6 to about 14 carbon atoms, $N(Q_1)(Q_2)$, OQ_1 , halo, SQ_1 or CN;

n is from 2 to 50; and

m is 0 or 1;

 R_2 is H, a hydroxyl protecting group, or an oligonucleotide; and

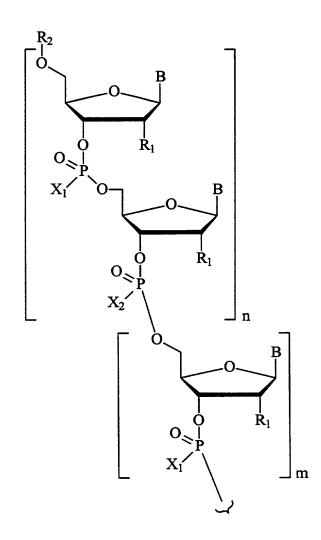
 $\ensuremath{R_3}$ is OH, an oligonucleotide, or a linker connected to a solid support.

- 8. The compound of claim 7 wherein R_1 is $-O-CH_2-CH_2-O-10$ CH_3 .
 - 9. The compound of claim 8 wherein R_2 is $H_{\mbox{\scriptsize ,}}$ and R_3 is OH.
- 10. The compound of claim 8 wherein R_2 is a phosphodiester-linked oligonucleotide or a phosphorothioate linked oligonucleotide.
 - 11. The compound of claim $8\ R_3$ is a phosphodiester-linked oligonucleotide or a phosphorothicate linked oligonucleotide.
- 12. R_2 and R_3 are each a phosphodiester-linked 20 oligonucleotide or a phosphorothioate linked oligonucleotide.
 - 13. A compound having the Formula:

(5') $W^1-W^2-W^3$ (3')

wherein:

 W^1 has the Formula:



wherein:

each B is a nucleobase;

one of X_1 or X_2 is O, and the other of X_1 or X_2 is S; each R_1 , is, independently, H, hydroxyl, C_1 - C_{20} alkyl, C_3 - C_{20} alkenyl, C_2 - C_{20} alkynyl, halogen, thiol, keto, carboxyl, nitro, nitroso, nitrile, trifluoromethyl, trifluoromethoxy, O-alkyl, S-alkyl, NH-alkyl, N-dialkyl, O-aryl, S-aryl, NH-aryl, O-aralkyl, S-aralkyl, NH-aralkyl, amino, N-

tides;

phthalimido, imidazole, azido, hydrazino, hydroxylamino, isocyanato, sulfoxide, sulfone, sulfide, disulfide, silyl, aryl, heterocycle, carbocycle, intercalator, reporter molecule, conjugate, polyamine, polyamide, polyalkylene 5 glycol, or polyether;

or R_1 is a group of formula $Z-R_{22}-(R_{23})_v$; $Z \text{ is O, S, NH, or } N-R_{22}-(R_{23})_v;$ $R_{22} \text{ is } C_1-C_{20} \text{ alkyl, } C_2-C_{20} \text{ alkenyl, or } C_2-C_{20}$ alkynyl;

10 R₂₃ is hydrogen, amino, halogen, hydroxyl, thiol, keto, carboxyl, nitro, nitroso, nitrile, trifluoromethyl, trifluoromethoxy, O-alkyl, S-alkyl, NH-alkyl, N-dialkyl, O-aryl, S-aryl, NH-aryl, O-aralkyl, S-aralkyl, NH-aralkyl, amino, N-phthalimido, imidazole, azido, hydrazino, hydroxylamino, isocyanato, sulfoxide, sulfone, sulfide, disulfide, silyl, aryl, heterocycle, carbocycle, intercalator, reporter molecule, conjugate, polyamine, polyamide, polyalkylene glycol, polyether, a group that enhances the pharmacodynamic properties of oligonucleotides, or a group that enhances the pharmacodynamic properties of oligonucleotides, or a group

v is from 0 to about 10; or R_1 has the formula:

$$-(O)_{y_1} - (CH_2)_{y_2} - O - N - (CH_2)_{y_2} - O - E$$

y1 is 0 or 1; y2 is independently 0 to 10; y3 is 1 to 10; $E \text{ is } C_1-C_{10} \text{ alkyl, } N\left(Q_1\right)\left(Q_2\right) \text{ or } N=C\left(Q_1\right)\left(Q_2\right);$

each Q_1 and Q_2 is, independently, H, C_1 - C_{10} alkyl, substituted alkyl, dialkylaminoalkyl, a nitrogen protecting group, a tethered or untethered conjugate group, a linker to a solid support; or Q_1 and Q_2 , together, are joined in a nitrogen protecting group or a ring structure that can include at least one additional heteroatom selected from N and O;

or R_1 has one of formula I or II:

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$$\begin{cases}
Z_0 - (CH_2)_{q1} & O \\
Z_1 & Z_3
\end{cases}$$

$$Z_1 = Z_3$$

$$Z_2 = Z_3$$

$$Z_3 = Z_3$$

$$Z_4 = Z_3$$

$$Z_4 = Z_3$$

$$Z_1 = Z_3$$

$$Z_2 = Z_3$$

$$Z_3 = Z_3$$

wherein

 Z_0 is O, S, or NH;

 q^1 is from 0 to 10;

 q^2 is from 1 to 10;

 q^3 is 0 or 1;

q4 is, 0, 1 or 2;

 Z_4 is OM_1 , SM_1 , or $N(M_1)_2$;

each M_1 is, independently, H, C_1-C_8 alkyl, C_1-C_8 20 haloalkyl, $C(=NH)N(H)M_2$, $C(=O)N(H)M_2$ or $OC(=O)N(H)M_2$;

 M_2 is H or C_1-C_8 alkyl;

 Z_1 , Z_2 and Z_3 comprise a ring system having from about 4 to about 7 carbon atoms, or having from about 3 to about 6 carbon atoms and 1 or 2 hetero atoms wherein said 25 hetero atoms are selected from oxygen, nitrogen and sulfur, and wherein said ring system is aliphatic, unsaturated aliphatic, aromatic, or saturated or unsaturated heterocyclic; and

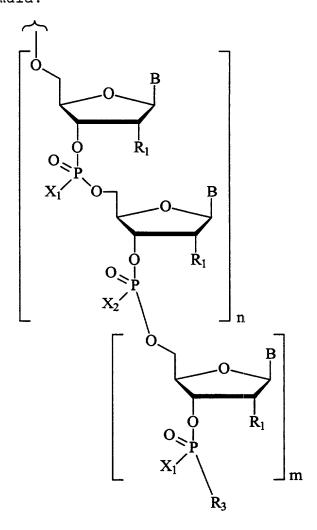
 Z_5 is alkyl or haloalkyl having 1 to about 10 carbon atoms, alkenyl having 2 to about 10 carbon atoms, alkynyl having 2 to about 10 carbon atoms, aryl having 6 to about 14 carbon atoms, $N\left(Q_1\right)\left(Q_2\right)$, OQ_1 , halo, SQ_1 or CN;

n is from 2 to 50; and

m is 0 or 1;

 $\ensuremath{\text{R}_2}$ is H, a hydroxyl protecting group, or an oligonucleotide;

 W^3 has the Formula:



10

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wherein R_3 is OH, an oligonucleotide, or a linker connected to a solid support; and

 \mathbf{W}^2 is a plurality of covalently bound nucleosides linked by phosphodiester or phosphorothicate linkages.

- 5 14. The compound of claim 13 wherein R_1 is $-O-CH_2-CH_2-O-CH_3$.
 - 15. The compound of claim 14 wherein R_2 is H, and R_3 is OH.
- 10 16. The compound of claim 14 wherein n is about 5 to about 50.
 - 17. The compound of claim 14 wherein n is about 8 to about 30.
- 18. The compound of claim 14 wherein n is about 4 to 15 about 15.
 - 19. The compound of claim 14 wherein n is 2 to about 10.
- 20. The compound of claim 14 wherein W^2 is a plurality of covalently bound nucleosides linked by phosphodiester 20 linkages.
 - 21. The compound of claim 14 wherein W^2 is a plurality of covalently bound nucleosides linked by phosphorothicate linkages.
- 22. A composition comprising a compound of claim 1 and 25 an acceptable carrier.

- 23. A composition comprising a compound of claim 7 and an acceptable carrier.
- 24. A composition comprising a compound of claim 12 and an acceptable carrier.
- 25. A method of modulating the production or activity of a protein in an organism, comprising contacting said organism with a compound of claim 1.
- 26. A method of modulating the production or activity of a protein in an organism, comprising contacting said 10 organism with a compound of claim 7.
 - 27. A method of modulating the production or activity of a protein in an organism, comprising contacting said organism with a compound of claim 13.

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- 28. A method of treating an organism having a disease characterized by the undesired production of a protein, contacting said organism with a compound of claim 1.
- 29. A method of treating an organism having a disease characterized by the undesired production of a protein,20 contacting said organism with a compound of claim 7.
 - 30. A method of treating an organism having a disease characterized by the undesired production of a protein, contacting said organism with a compound of claim 13.
- 31. A method of assaying a nucleic acid, comprising 25 contacting a solution suspected to contain said nucleic acid with a compound of claim 1.

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32. A method of assaying a nucleic acid, comprising contacting a solution suspected to contain said nucleic acid with a compound of claim 7.

33. A method of assaying a nucleic acid, comprising5 contacting a solution suspected to contain said nucleic acid with a compound of claim 13.